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(54) Coating for bioprosthetic device and method of making same.

(57) A coating and integral treatment for improving the biophysical stability of bioprosthetic devices after implantation, particularly with respect to calcification, and, specifically, a method for treating animal tissues, such as heart valves, to provide improved biophysical stability in allograft and heterograft transplants. The increased stability results, in-part, from the creation of a three-dimensional matrix of a primary structural component of the prosthetic device and covalently attached calcification inhibitors. Other materials, some having additional stabilizing effects, may be utilized to form additional bridges or fill the interstitial gaps in the matrix. After implantation, the resultant modified device exhibits minimal surface for intramatrix growth of calcium phosphate crystals, and additionally, may inhibit platelet aggregation, enzymatic degradation and host rejection, while minimizing the risk of mechanical failure, in the host organism.

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COATING FOR BIOPROSTHETIC DEVICEANDMETHOD OF MAKING SAMEBACKGROUND OF THE INVENTION

For many years, a variety of animal tissues, as well as some synthetic polymers, have been used to make prosthetic devices for surgical implantation into human beings and other animals. However, because these devices are different on a molecular level from the host organism's own tissue, they usually elicit a wide variety of reactions in the host. The response is manifested by a low-grade, rapid deterioration of the transplant, which in turn, mandates additional surgery.

To improve the longevity of transplanted devices, a number of remedies have been proposed. In the processing of natural tissues, a common stabilization technique involves treatment with tanning agents, such as formaldehyde. Glutaraldehyde, a well known cross-linking agent, has also been used with success in this regard. In fact, a number of studies have shown that heart valves treated with glutaraldehyde can remain functional in situ for many years. However, recent research has indicated that such glutaraldehyde preserved implantations can still elicit significant host reactions, including calcification, fibrin deposition and an anaphylactic response. (For example, see Slanczka, D.J. and Bajpai, P.K., "Immunogenicity of Glutaraldehyde-treated Porcine Heart Valves", IRCS Medical Science: Bio-Technology; Cardiovascular System; Immunology and Allergy; Pathology; Surgery and Transplantation;

6, 421 (1978).)

It has also been theorized that natural prosthetics may be biodegradable, and thus labile even after short placement periods. In vitro enzyme degradation of the tissues prior to implantation has been utilized to minimize this obstacle, but this degradation is not totally effective in mitigating the antigenic response; and moreover, the tissue can lose significant portions of its inherent structural framework, which can cause further mechanical weakening of the entire device.

Although considerable success has been achieved by implanting synthetic devices instead of natural devices, at present, they also present significant difficulties. There is a substantial biological failure rate among these devices due to incompatibility with biological tissues. After removal of the implant, fibrin layering, aneurysm formation, lipid deposition and many clinical malfunctions have been noted.

A further problem, common to many of the synthetic and natural prosthetics alike, is minimal flexibility. Glutaraldehyde-treated natural devices are often cross-linked to such a degree that much of their natural flexibility is lost, and after prolonged periods of implantation, brittleness often becomes even more pronounced. Similarly, synthetic devices generally become increasingly hardened after prolonged implantation.

Therefore, there is a recognized need for an improved treatment of prosthetic devices prior to implantation, which will render these devices more

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durable, yet minimize negative host responses. The present invention fulfills this need.

#### SUMMARY OF THE INVENTION

In accordance with one aspect of the invention, a coating for heart valves and other prosthetic devices is provided that has greatly improved biophysical stability after the device is implanted in a host organism. Through the formation of a three-dimensional cross-linked matrix primarily composed of a calcification inhibitor covalently bound to accessible regions of the device, a substantially non-antigenic bioprosthesis with minimum calcification sites may be produced.

Suitable calcification inhibitors include natural protein polysaccharides, such as chondroitin sulfates and hyaluronate. Generally, sulfated polysaccharides are preferred, but diphosphonates, phosphoproteins, dyes, such as alzarin red S and methylene blue, and other polyanions may be used.

The incorporation of other agents into the matrix can further enhance long term survival of the implanted device. Specifically, bridging agents, such as diamines, that create additional cross-linking sites and additional covalent binding sites for attaching other specified materials, such as antithrombogens, may be bound to the matrix. Also, the presence of materials that fill the interstitial gaps in the matrix can provide greater stability by limiting nucleation and the growth of hydroxyapatite crystals.

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Another aspect of the invention is a process for treating bioprosthesis to provide a coating, such as described above, for improved stability after implantation. The method, which can utilize the compounds described above, comprises the steps of:

5 harvesting tissue from an organism; initiating a number of covalent cross-links, preferably with glutaraldehyde, in the protein structure of the tissue sufficient to protect the tissue from initial

10 losses in structural integrity; soaking the tissue in calcification inhibitor; covalently binding the calcification inhibitor to the tissue, preferably with a carbodiimide; and sterilizing. Additional steps may include the covalent binding of bridging agents,

15 such as diamines, antithrombogenic agents and gap filling materials to the tissue. The treatment is particularly useful for rendering animal connective tissues, such as mammalian heart valves and blood vessels, substantially water insoluble and less likely

20 to initiate calcification than natural tissue or tanned tissue.

#### DETAILED DESCRIPTION OF THE INVENTION

##### AND

#### PREFERRED EMBODIMENTS

Exemplary starting materials useful in practicing the invention include: animal tissues of diverse origin, e.g. heart valves, blood vessels,

25 pericardium, dura mater, ligaments, tendons, and other collagen-rich tissues, as well as reconstituted or native collagen fibers and other materials with accessible cross-linking sites. Assuming tissues are used, they are first cleaned from adherent fat or

30 loose connective tissue as soon as possible after

harvesting. Immediately thereafter, they are placed in a balanced electrolyte solution that is calcium-free and buffered at a neutral pH with a phosphate buffer. This solution, kept cool (4-8 degrees Centigrade), contains a calcium chelator, such as EDTA-Na at about 0.05 molar concentration, to sequester calcium present in the tissue.

The following steps are then utilized to adequately cross-link and modify the tissue in this exemplary process:

- 1) Immediately after harvesting and cleaning, the tissue is placed in a solution containing 0.05 wt. % glutaraldehyde buffered with phosphate at pH 7.0, and made isotonic with a calcium free, balanced electrolyte solution. This causes partial cross-linking of the collagen and the protein-like compounds naturally associated with it (called protein-polysaccharides) and is performed to prevent swelling and distortion of the ultrastructure of the connective tissue.
- 2) The tissue is then placed in a solution containing a calcification inhibitor, preferably chondroitin sulfate at a concentration of about 0.5 to about 5 wt. %, preferably about 1.0 wt. %. Chondroitin sulfate is available commercially or may be prepared from a variety of cartilagenous sources. In some instances, it may be desirable to use the protein-polysaccharides associated with collagen in natural tissues.

5 These include chondroitin-6-sulfate,  
chondroitin-4-sulfate and hyaluronate.  
Generally, polysaccharides of the  
chondroitin sulfate variety that are  
rich in weak negative charges (carboxyl  
groups) and in strong negative charges  
(sulfate groups), such as sulfated  
polysaccharides, are preferred. Other  
substances that are known inhibitors of  
10 calcification include diphosphonates,  
which are characterized by the presence  
of a P-C-P or a P-N-P bond. It is  
theorized that P-C-P and P-N-P bonds are  
not "bio-degradable" and are, therefore,  
15 very stable in tissues. A typical  
diphosphonate is 3-amino-1-hydroxy-  
propane 1, 1-diphosphonic acid. Other  
diphosphonates with active amino or  
carboxyl groups can easily be attached  
20 by covalent bonds and act as inhibitors  
of calcification at the surface or with-  
in the interstitial spaces of matrices  
formed. Additional calcification  
inhibitors include phosvitin or other  
25 phosphoproteins, dyes, such as alizarin  
red S, and methylene blue, calcium  
chelators, such as EDTA and EGTA, and  
other polyanions. The calcium inhibitor  
chosen is preferably allowed to diffuse  
30 freely into the tissue, usually until  
equilibrium is reached, which is after  
about 12 hours.

- 3) To the solution containing the calcium inhibitor and the tissue, an aliphatic

diamine, preferably hexanediamine, is added to provide additional binding sites and cross-links in the subsequent covalent binding steps. Although diamines are preferable, other compounds with free terminal amino or carboxyl groups can be utilized. The diamine and chondroitin sulfate may be added to the solution at the same time, but by adding the calcium inhibitor first, more polyelectrolytes are probably allowed to diffuse into the tissue.

- 4) The tissues and additives are then cross-linked by a water-soluble carbodiimide. Carbodiimides apparently form peptide bonds by activation of carboxyl groups to allow reaction with amino groups. The cross-linking occurs at a carbodiimide concentration of about 0.02 to about 0.1 molar, preferably about 0.05 molar, in a balanced electrolyte solution. The pH should be between about 4.7 and about 5.2, and is maintained at about 5.0 by the addition of HCl. The preferred carbodiimide is 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide HCl. If desired, ethanol and other organic solvents may be added to decrease the dielectric constant. The cross-linking reaction is allowed to proceed from about 30 minutes up to 10 hours or more.

- 5) After coupling is completed, the excess reagents are removed by washing with a



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balanced electrolyte solution at a neutral pH, which also contains 0.05 M EDTA.

5                   6) The tissue is then transferred to a neutral pH buffered solution, containing about 0.2 to about 0.5 wt. % glutaraldehyde, preferably about 0.3 wt. %, in a balanced electrolyte environment. This  
10 final solution can be supplemented with alcohol at a concentration of about 20 to about 50 wt. %, and surfactants, such as anionic alkyl sulfates or alcohol and Formaldehyde, for sterility and storage.

15                   It is possible to modify the procedure stated above. For example, repeating the equilibration with the calcification inhibitor and subsequently reactivating the entire matrix with carbodiimide may be desirable.

20                   Moreover, prior to or in conjunction with the final glutaraldehyde treatment, anti-thrombogenic compounds, such as heparin, which may also convey additional attributes, may be added at a concentration of about 0.2 to about 1.0 wt. %. Also, globular  
25 proteins, small molecular weight peptides, or poly-electrolytes, such as polylysine or polyglutamic acid, or mixed copolymers of poly-electrolytes, may be added. By allowing these materials to diffuse into the cross-linked matrix, further bridging between the  
30 tissue components and exogenous materials may occur, and the interstitial gaps may be filled. It is believed that by filling the interstitial gaps the deposit of calcium ions is minimized and hydroxyapatite and other crystal growth may be substantially

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inhibited.

Some basic improvements provided by the present invention will now be discussed.

#### I. Immunogenicity

5        If material implanted in an organism can be rendered insoluble, antigenicity can be substantially eliminated. To be recognized, antigens must be presented in a soluble form to activate the immune system of the host organism. In many cases, materials that  
10        are insoluble at the time of implantation can be rendered soluble by naturally occurring enzymatic or chemical processes. It is believed that the introduction of sufficient cross-links prohibits the enzyme systems of the host from solubilizing the implanted  
15        material, thereby essentially eliminating antigenicity.

      Glutaraldehyde treatment also introduces cross-links, but for reasons not completely understood, the cross-links generated in the present  
20        invention render the entire device even less soluble. Without being bound to any particular reason, perhaps this reduced solubility is due to the presence of cross-links different than those created when glutaraldehyde is used alone. Since glutaraldehyde  
25        apparently acts primarily on lysine residues, the type and amount of bridges produced are somewhat limited. The present invention enhances the amount of cross-linking by covalently attaching new amino groups to the structure, and additionally allows the use of  
30        other moieties, such as peptide bound glutamic and aspartic acids, to attach more cross-links in different locations by the carbodiimide reaction

mechanism.

## II. Calcification

A significant, but often ignored, problem associated with the implantation of grafts rich in collagen and elastin is the propensity of these grafts to induce calcification. Collagen in particular has an intrinsic ability to calcify, and a mixture of collagen fibers with saturated solutions of calcium and phosphate ions will induce nucleation, which is closely followed by crystal growth. The addition of polyanions, particularly sulfated polysaccharides, can essentially prevent this nucleation process.

Some sulfated polysaccharides, such as endogenous chondroitin sulfate, can be bound to the collagen during the tanning procedure. But, these polyanions are usually degraded by the host and subsequently removed from the graft. Therefore, the initial protection afforded to the tissue by these materials is lost, and exposure of the functional groups in collagen, as well as the new open spaces generated, can now greatly enhance nucleation of calcium and phosphate ions. The process used in the present invention covalently links these polyanions to collagen, or some other primary structural component of the prosthetic device, and sufficiently cross-links the entire structure to prevent degradation and crystal growth. The addition of any extraneous calcification inhibitors that are also bound and cross-linked can further minimize calcification.

## III. Host Induced Graft Destruction

Uncross-linked implanted, fresh heterografts

or allografts are rapidly destroyed by the defense mechanisms of a host organism. Adequate cross-linking, which as previously discussed insolubilizes the tissue, can prevent this destruction. Again, although glutaraldehyde induces a certain number of cross-links, these have been shown to be inadequate. Apparently, because of the different nature of the cross-links produced in the present invention, greater stability can be obtained, while the actual density of cross-links may be fewer. This is possible because the cross-links have been designed to span a broader set of distances, both inter- and intra-molecular; as well as to join not only lysine residues present, but also the free carboxyl groups of glutamic and aspartic acid. Apparently, these different types of cross-links give added resistance to the treated tissues against enzymatic degradation, but importantly, without significant decreases in the mechanical attributes of the grafts.

#### IV. Compatibility With Blood Surfaces

Collagen, the primary structural component of most animal tissues, is a well known platelet aggregator and blood clot initiator. Since the connective tissues used in prostheses are very rich in collagen, the present invention utilizes substances capable of reducing the tissues' thrombogenic potential. Chondroitin sulfate also serves this purpose, but additional compounds with antithrombogenic properties, such as heparin, may be used. These compounds, once covalently bound, substantially decrease the ability of collagen to aggregate platelets, thereby significantly decreasing the probability of thrombus formation.

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## V. Changes in Mechanical Properties

The function of a transplanted device under most circumstances will depend on the retention of adequate visco-elastic behavior at a level particularly suitable for the function that the graft is to perform. Maintaining the proper amount of elasticity depends in-part on the degree of cross-linking. Insufficient cross-links could allow for flow, enzymatic degradation, and subsequent destruction of the physical integrity of the device. On the other hand, too many cross-links can be conducive to brittleness, and result in loss of function. The present invention provides an adequate number of cross-links to help retain the structural integrity of the implanted device, but not so many or so clustered that elasticity is lost.

ALTERNATIVE EMBODIMENT

Tissues are received from the slaughter house, cleaned to remove loosely adhering material, and rinsed with cold phosphate-buffered, physiological saline.

The tissues are then processed, usually at about 4 degrees Centigrade, as follows:

- 1) treat with a glutaraldehyde solution at a concentration between about 0.05 wt. % and about 0.4 wt. %, preferably about 0.15 wt. %, for between about 12 and about 64 hours, preferably about 48 hours;
- 2) rinse the tissue in phosphate buffered

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saline to remove non-reacted glutaraldehyde;

- 5 3) place the tissue in a solution with a pH of about 7.4 containing between about 0.1 wt. % and about 2.0 wt. % hexanediamine, preferably about 0.5 wt. %;
- 4) incubate for about 2 to about 10 hours, preferably 4 hours;
- 10 5) transfer the tissues to a buffered saline solution at a pH of about 4.9 that contains between about 0.1 and about 1.0 wt. % of a water soluble carbodiimide, preferably 0.5 wt. % 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide HCl;
- 15 6) incubate for about 30 minutes to about 10 hours, preferably 1 hour, while maintaining the pH of the entire mixture at about pH 5.0 with an aqueous HCl solution;
- 20 7) place the tissue in neutral, phosphate buffered saline for rinsing from about 2 to about 12 hours, preferably about 6 hours;
- 25 8) place the tissue in a buffered neutral saline solution that contains between about 0.5 and about 3 wt. % of a sulphated polysaccharide, preferably about 1.0 wt. % chondroitin sulfate from
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whale and shark cartilage, the sodium salt of mixed isomers (No. C-3129, Sigma Chemical Company);

- 5 9) incubate for about 6 to about 16 hours, preferably 12 hours, until equilibration (gentle mechanical shaking may be used);
- 10 10) transfer the tissue to a buffered saline solution, at a pH of about 4.9 that contains about 0.1 to about 5 wt. % of a water soluble carbodiimide, preferably about 0.5 wt. % of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide HCl, and about 0.1 to about 5 wt. % of an aliphatic diamine, preferably about 15 0.5 wt. % hexanediamine;
- 20 11) incubate for about 30 minutes to about 10 hours, preferably 1 hour, while maintaining the pH of the entire mixture at about pH 5.0 with an aqueous HCl solution;
- 25 12) rinse the tissue in a neutral, phosphate buffered saline solution from about 2 to about 12 hours, preferably about 6 hours;
- 30 13) transfer the tissue to a neutral, phosphate buffered saline solution containing between about 0.2 and about 2.0 wt. % of an antithrombogenic agent, preferably 1.0 wt. % heparin;

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- 14) incubate for about 30 minutes to about 10 hours, preferably about 1 hour;
- 15) add, to the solution, glutaraldehyde to a final concentration of between about 0.1 and about 1.0 wt. %, preferably about 0.4 wt. %;
- 16) incubate for about 30 minutes to about 10 hours, preferably about 1 hour;
- 17) transfer to a final storage, neutral, phosphate buffered solution containing about 0.4 wt. % glutaraldehyde, preferably about 0.4 wt. %, about 0.2 to about 2.0 wt. % of formaldehyde, preferably about 1.0 wt. % and about 20 to about 40 wt. % alcohol, preferably about 30 wt. %.

Although the invention has been described in detail, it will be understood by one of ordinary skill in the art that various modifications can be made without departing from the spirit and scope of the invention. Accordingly, it is not intended that the invention be limited except as by the appended claims.



CLAIMS

1. A process for improving the biophysical stability of bioprosthesis for hetrograft or allograft implantation, which comprises;

Harvesting tissue from an organism; initiating covalent cross-links in the protein structure of the tissue to protect the tissue from excessive swelling or other losses of structural integrity, preferably by reacting the tissue with Glutaraldehyde; soaking the tissue in an aqueous solution of a calcification inhibitor; covalently binding the calcification inhibitor to the tissue thereby forming a three dimensional matrix, for example, by reacting the tissue with a carbodiimide, preferably 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide; and sterilizing the matrix, preferably in a solution containing formaldehyde or alcohol or a mixture thereof.

2. A process according to Claim 1, wherein the calcification inhibitor is a polyanion, preferably an anionic polysaccharide.

3. A process according to Claim 1, wherein the calcification inhibitor is a sulfated polysaccharide, preferably chondroitin-4-sulfate, chondroitin-6-sulfate, hyaluronate or a mixture of any two or more thereof.

4. A process according to any one of Claims 1 to 3, wherein the tissue is treated prior to or simultaneously with the covalent binding step to the action of a bridging agent, for example, a diamine, preferably an aliphatic diamine, that can covalently bind to the tissue thereby providing additional sites for peptide bond formation.

5. A process according to any one of Claims 1 to 4, which further includes the step of covalently bonding an antithrombogenic agent, preferably heparin to the matrix.

6. A process according to any one of Claims 1 to 5, wherein the tissue is an animal connective tissue, preferably a mammalian heart valve, blood vessel, pericardium, dura mater, ligament, tendon or other collagen-rich tissue.

7. A coating for a prosthetic device that provides increased stability for allograft or heterograft implantations, said coating comprising:

a three dimensional, cross-linked matrix of an exogenous calcification inhibitor covalently bound to accessible regions of the device wherein the coating is substantially non-antigenic and has minimal calcification initiation sites.

8. A process for treating heart valves prior to implantation into a human comprising the steps of:

harvesting a fresh heart valve from a donor organism;

initiating cross-links in the valve by treating with glutaraldehyde;

incubating the valve in a solution containing a diamine;

reacting the valve with a water soluble carbodiimide;

soaking the valve in a solution containing a sulphated polysaccharide;

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reacting the valve with a water soluble carbodiimide in the presence of a diamine; soaking the valve in a solution containing heparin;

reacting the valve with glutaraldehyde; storing the valve in a sterilizing solution.

9. A process for improving the biophysical stability of bioprotheses for heterograft or allograft implantation, which comprises:

harvesting tissue from an organism; initiating covalent cross-links in the protein structure of the tissue to protect the tissue from excessive swelling or other losses of structural integrity; soaking the tissue in an aqueous solution of a diphosphonate;

covalently binding the diphosphonate to the tissue thereby forming a three-dimensional matrix; and sterilizing the matrix;

wherein the modified tissue produced is substantially water insoluble; and, after implantation in a host organism, the matrix is less likely to elicit an antigenic response or to be subject to calcification than natural tissue or tanned tissue.

10. A process for improving the biophysical stability of bioprostheses for heterograft or allograft implantation, which comprises:

harvesting tissue from an organism;  
initiating covalent cross-links in the protein structure of the tissue to protect the tissue from excessive swelling or other losses of structural integrity;  
soaking the tissue in an aqueous solution of a dye;  
covalently binding the dye to the tissue thereby forming a three-dimensional matrix; and  
sterilizing the matrix;  
wherein the modified tissue produced is substantially water insoluble; and, after implantation in a host organism, the matrix is less likely to elicit an antigenic response or to be subject to calcification than natural tissue or tanned tissue.

11. A process for improving the biophysical stability of bioprostheses for heterograft or allograft implantation, which comprises:
- harvesting tissue from an organism;
  - initiating covalent cross-links in the protein structure of the tissue to protect the tissue from excessive swelling or other losses of structural integrity;
  - soaking the tissue in an aqueous solution of a phosphoprotein;
  - covalently binding the phosphoprotein to the tissue thereby forming a three-dimensional matrix; and
  - sterilizing the matrix;

wherein the modified tissue produced is substantially water insoluble; and, after implantation in a host organism, the matrix is less likely to elicit an antigenic response or to be subject to calcification than natural tissue or tanned tissue.

12. A process for improving the biophysical stability of bioprostheses for heterograft or allograft implantation, which comprises:

harvesting the tissue from an organism;

initiating covalent cross-links in the protein structure of the tissue to protect the tissue from excessive swelling or other losses of structural integrity;

soaking the tissue in an aqueous solution of a chelating agent;

covalently binding the chelating agent to the tissue thereby forming a three-dimensional matrix; and,

sterilizing the matrix;

wherein the modified tissue produced is substantially water insoluble; and, after implantation in a host organism, the matrix is less likely to elicit an antigenic response or to be subject to calcification than natural tissue or tanned tissue.

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